Reply Dated April 21, 2008 After Final Office Action of December 21, 2007

AMENDMENTS TO THE CLAIMS

1. (Currently Amended) A method for preparing a expanding cytotoxic lymphocyte

lymphocytes which comprises:

expanding a cytotoxic lymphocyte culturing peripheral blood mononuclear cells, Natural

Killer (NK) cells, umbilical cord blood mononuclear cells, hematopoietic stem cells or blood

components containing these cells in the presence of [[a]] at least one fibronectin fragment or a

mixture thereof together with interleukin-2,

wherein the fibronectin fragment is

[[i)]] a polypeptide comprising at least one of the amino acid sequences of an amino acid

sequence selected from the group consisting of SEO ID NOS: 1 to 19, [[orl]]

ii) a polypeptide having a substitution of one or more amino acids in the amino acid

sequence of the polypeptide of i), and having a function which is equivalent to that of the

polypeptide of i), wherein the substitution of one or more amino acids is a substitution within

each of the groups of:

a) glycine, alanine:

b) valine, isoleucine, leucine;

c) aspartic acid, glutamic acid, asparagine, glutamine;

d) serine, threonine;

e) lysine, arginine; and

f) phenylalanine, tyrosine

wherein said culturing is performed for 2-15 days.

2.

MAA/TJS/mrh

Docket No.: 1422-0644PUS1

After Final Office Action of December 21, 2007

2. (Currently Amended) The method according to claim 1, wherein the prepared

expanded cytotoxic lymphocytes highly expresses express an interleukin-2 receptor

at a higher level than [[a]] cytotoxic lymphocyte prepared lymphocytes expanded in the absence

of [[a]] at least one fibronectin fragment or a mixture thereof.

3. (Currently Amended) The method according to claim 1, wherein the prepared

expanded cytotoxic lymphocyte-expresses lymphocytes express more CD8 than [[a]] cytotoxic

lymphocyte prepared lymphocytes expanded in the absence of [[a]] at least one fibronectin

fragment or a mixture thereof.

4. (Currently Amended) The method according to any one of claims 1 to 3, wherein the

prepared expanded cytotoxic lymphocyte maintains lymphocytes maintain cytotoxic activity

longer than [[a]] cytotoxic lymphocyte prepared lymphocytes expanded in the absence of [[a]] at

least one fibronectin fragment or a mixture thereof.

5. (Currently Amended) The method according to claim 1, wherein said at least one

fibronectin fragment or a mixture thereof is immobilized on a solid phase.

6. (Previously Presented) The method according to claim 5, wherein the solid phase is a

cell culture vessel or a cell culture carrier.

3

Application No. 10/509,055 Docket No.: 1422-0644PUS1 Reply Dated April 21, 2008

After Final Office Action of December 21, 2007

7. (Previously Presented) The method according to claim 6, wherein the cell culture

vessel is a petri dish, a flask or a bag, and the cell culture carrier is beads, a membrane or a slide

glass.

8. (Withdrawn) The method according to claim 1, wherein expanding a cytotoxic

lymphocyte is performed in a cell culture medium comprising said fibronectin fragment or a

mixture thereof.

9. (Cancelled)

10. (Currently Amended) The method according to claim 1, wherein the at least one

fibronectin fragment has cell adhesion activity and/or heparin binding activity.

11. (Cancelled)

12. (Currently Amended) The method according to claim 1, comprising:

expanding a cytotoxic lymphocyte in a cell culture in the presence of said at least one

fibronectin fragment or a mixture thereof,

wherein at least (a) or (b) is true:

(a) a ratio of the number of cells present at the initiation of the cell culture to a cell

4

culture area is 1 cell/cm2 to 5 × 105 cells/cm2; and

Application No. 10/509,055

Reply Dated April 21, 2008

Docket No.: 1422-0644PUS1

After Final Office Action of December 21, 2007

(b) a concentration of cells at the initiation of the cell culture is from 1 cell/ml to $5 \times$

105 cells/ml.

13. (Cancelled)

14. (Withdrawn) A cytotoxic lymphocyte obtained by the method of claim 1.

15. (Withdrawn) A medicament comprising as an effective ingredient a cytotoxic

lymphocyte obtained by the method of claim 1.

16. (Withdrawn) An agent for enhancing an interleukin-2 receptor expression of a cell,

characterized in that the agent comprises as an effective ingredient fibronectin, a fragment

thereof or a mixture thereof.

17. (Withdrawn) The agent according to claim 16, wherein the fibronectin fragment is a

polypeptide comprising at least one of the amino acid sequences represented by SEQ ID NOs: 1

to 7 of Sequence Listing, or a polypeptide having substitution, deletion, insertion or addition of

one or more amino acids in the amino acid sequence of said polypeptide, wherein the

polypeptide has functions equivalent to that of said polypeptide.

18. (Withdrawn) The agent according to claim 17, wherein the fibronectin fragment has

5

cell adhesion activity and/or heparin binding activity.

19. (Withdrawn) The agent according to claim 17, wherein the fibronectin fragment is a

polypeptide selected from polypeptides comprising any one of the amino acid sequences shown

in SEO ID NOs: 8 to 19 of Sequence Listing.

20. (Withdrawn) An agent for improving a ratio of CD8-positive cell in a lymphocyte,

characterized in that the agent comprises as an effective ingredient fibronectin, a fragment

thereof or a mixture thereof.

21. (Withdrawn) The agent according to claim 20, wherein the fibronectin fragment is a

polypeptide comprising at least one of the amino acid sequences represented by SEQ ID NOs: 1

to 7 of Sequence Listing, or a polypeptide having substitution, deletion, insertion or addition of

one or more amino acids in the amino acid sequence of said polypeptide, wherein the

polypeptide has functions equivalent to that of said polypeptide.

22. (Withdrawn) The agent according to claim 21, wherein the fibronectin fragment has

cell adhesion activity and/or heparin binding activity.

23. (Withdrawn) The agent according to claim 21, wherein the fibronectin fragment is a

polypeptide selected from polypeptides comprising any one of the amino acid sequences shown

6

in SEQ ID NOs: 8 to 19 of Sequence Listing.

MAA/TJS/mrh

Docket No.: 1422-0644PUS1

Docket No.: 1422-0644PUS1 Application No. 10/509,055

Reply Dated April 21, 2008 After Final Office Action of December 21, 2007

24. (Withdrawn) An agent for improving or maintaining cytotoxic activity in a cytotoxic

lymphocyte, characterized in that the agent comprises as an effective ingredient fibronectin, a

fragment thereof or a mixture thereof.

25. (Withdrawn) The agent according to claim 24, wherein the fibronectin fragment is a

polypeptide comprising at least one of the amino acid sequences represented by SEQ ID NOs: 1

to 7 of Sequence Listing, or a polypeptide having substitution, deletion, insertion or addition of

one or more amino acids in the amino acid sequence of said polypeptide, wherein the

polypeptide has functions equivalent to that of said polypeptide.

26. (Withdrawn) The agent according to claim 25, wherein the fibronectin fragment has

cell adhesion activity and/or heparin binding activity.

27. (Withdrawn) The agent according to claim 25, wherein the fibronectin fragment is a

polypeptide selected from polypeptides comprising any one of the amino acid sequences shown

in SEO ID NOs: 8 to 19 of Sequence Listing.

28. (Currently Amended) A method for increasing expression of an interleukin-2 receptor

in [[al] cytotoxic lymphocyte lymphocytes, which comprises:

expanding a cytotoxic lymphocyte culturing peripheral blood mononuclear cells, Natural

Killer (NK) cells, umbilical cord blood mononuclear cells, hematopoietic stem cells or blood

components containing these cells in the presence of [[all at least one fibronectin fragment or a 7

After Final Office Action of December 21, 2007

mixture thereof together with interleukin-2, thereby increasing expression of

interleukin-2 receptor in a cytotoxic lymphocyte the cells,

wherein the fibronectin fragment is

[[i]] a polypeptide comprising at least one of the amino acid sequences of an amino acid

sequence selected from the group consisting of SEQ ID NOS: 1 to 19, [[or]]

ii) a polypeptide having a substitution of one or more amino acids in the amino acid

sequence of the polypeptide of i), and having a function which is equivalent to that of the

polypeptide of i), wherein the substitution of one or more amino acids is a substitution within

each of the groups of:

a) glycine, alanine;

b) valine, isoleucine, leucine:

c) aspartic acid, glutamic acid, asparagine, glutamine;

d) serine, threonine:

e) lysine, arginine; and

f) phenylalanine, tyrosine

wherein said culturing is performed for 2-15 days.

29. (Currently Amended) A method for increasing the number of CD8-positive cells in a

cytotoxic lymphocyte population lymphocytes, which comprises:

expanding a cytotoxic lymphocyte culturing peripheral blood mononuclear cells, Natural

Killer (NK) cells, umbilical cord blood mononuclear cells, hematopoietic stem cells or blood

components containing these cells in the presence of [[a]] at least one fibronectin fragment or a 8

After Final Office Action of December 21, 2007

mixture thereof together with interleukin-2, thereby increasing the number of [[CD-8]] CD8-

Docket No.: 1422-0644PUS1

positive cells in a cytotoxic lymphocyte population the cultured cells,

wherein the fibronectin fragment is

[[i]] a polypeptide comprising at least one of the an amino acid sequences sequence

selected from the group consisting of SEO ID NOS: 1 to 19, [[or]]

ii) a polypeptide having a substitution of one or more amino acids in the amino acid

sequence of the polypeptide of i), and having a function which is equivalent to that of the

polypeptide of i), wherein the substitution of one or more amino acids is a substitution within

each of the groups of:

a) glycine, alanine;

b) valine, isoleucine, leucine;

c) aspartic acid, glutamic acid, asparagine, glutamine;

d) serine, threonine;

e) lysine, arginine; and

f) phenylalanine, tyrosine

wherein said culturing is performed for 2-15 days.

30. (Currently Amended) A method for improving or maintaining cytotoxic activity in

[[a]] cytotoxic lymphocyte lymphocytes, which comprises:

expanding a cytotoxic lymphocyte culturing peripheral blood mononuclear cells, Natural

Killer (NK) cells, umbilical cord blood mononuclear cells, hematopoietic stem cells or blood

components containing these cells in the presence of [[a]] at least one fibronectin fragment or a 9

MAA/T.IS/mrh

Docket No.: 1422-0644PUS1

mixture thereof together with interleukin-2, thereby improving or maintaining cytotoxic activity in a cytotoxic lymphocyte the cells,

wherein the fibronectin fragment is

[[i)]]a polypeptide comprising at least one of the an amino acid sequences sequence

selected from the group consisting of SEO ID NOS: 1 to 19, [[or]]

ii) a polypeptide having a substitution of one or more amino acids in the amino acid

sequence of the polypeptide of i), and having a function which is equivalent to that of the

polypeptide of i), wherein the substitution of one or more amino acids is a substitution within

each of the groups of:

a) glycine, alanine;

b) valine, isoleucine, leucine:

e) aspartic acid, glutamic acid, asparagine, glutamine;

d) serine, threonine;

e) lysine, arginine; and

f) phenylalanine, tyrosine

wherein said culturing is performed for 2-15 days.

31. (Currently Amended) The method according to claim 1, further comprising

transducing a foreign gene into [[a]] the cytotoxic lymphocytes.

32. (Original) The method according to claim 31, wherein the foreign gene is transduced

10

using retrovirus, adenovirus, adeno-associated virus or simian virus.

Application No. 10/509,055
Reply Dated April 21, 2008

Docket No.: 1422-0644PUS1

After Final Office Action of December 21, 2007

33. (Currently Amended) The method according to claim 1, wherein an expansion ratio

of the cytotoxic lymphocyte lymphocytes is high as compared to that of [[the]] a method for

preparing expanding [[a]] cytotoxic lymphocyte lymphocytes in the absence of [[a]] at least one

fibronectin fragment or a mixture thereof.

34. (Currently Amended) The method according to claim 1, wherein expanding [[a]]

cytotoxic lymphocyte lymphocytes is performed in the presence of both of said at least one

fibronectin fragment or mixture thereof and an anti-CD3 antibody.

35. (Currently Amended) The method according to claim 1, wherein expanding [[a]]

cytotoxic lymphocyte <u>lymphocytes</u> is performed by incubating peripheral blood mononuclear

11

cells or umbilical cord blood mononuclear cells.

36. (Cancelled)